

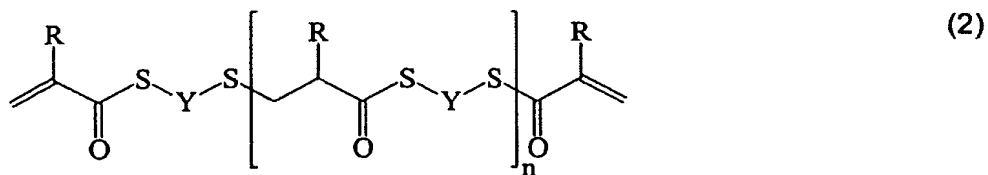
Mixtures for preparing transparent plastics,
transparent plastics, and process for their
preparation, and use

5 The present invention relates to mixtures for preparing
transparent plastics. The present invention further
relates to transparent plastics which can be prepared
from the mixtures, and to a process for their
preparation. The present invention also relates to the
10 use of transparent plastics for producing optical,
especially ophthalmic, lenses.

Spectacles have become an essential component of
everyday life. Among these, spectacles with plastics
15 lenses have in particular gained importance recently,
because they weigh less and are less breakable than
spectacle lenses composed of inorganic materials, and
can be coloured by means of suitable dyes. The
production of plastics spectacle lenses generally uses
20 high-transparency plastics obtainable, by way of
example, from diethylene glycol bis(allyl carbonate)
(DAC), thiourethane compounds having α,ω -terminated
multiple bonds or sulphur-containing (meth)acrylates.

25 DAC plastic has very good impact strength and
transparency, and good processibility. However, a
disadvantage is that the relatively low refractive
index n_D of about 1.50 requires that both the centre and
the edges of these plastics lenses be reinforced, the
30 spectacle lenses being correspondingly thick and heavy.
This markedly reduces the wearer comfort of spectacles
with DAC plastics lenses.

The specification DE 4234251 discloses sulphur-
35 containing polymethacrylates which are obtained via
free-radical copolymerization of a monomer mixture
composed of compounds of the formula (1) and (2)



Here, Y is an unbranched or branched, acyclic or cyclic alkyl radical having from 2 to 12 carbon atoms, or an aryl radical having from 6 to 14 carbon atoms, or an alkaryl radical having from 7 to 20 carbon atoms, and the carbon chains here may have interruption by one or more ether or thioether groups. R is hydrogen or methyl and n is a whole number in the range from 1 to 6.

10

According to DE 4234251, the monomers of the formula (1) and (2) generally have a molar ratio of from 1:0.5 to 0.5:1. The monomer mixture is prepared via reaction of at least two mols of (meth)acryloyl chloride or (meth)acrylic anhydride with one mole of a dithiol, by reacting the (meth)acryloyl chloride or (meth)acrylic anhydride in an inert organic solvent and the dithiol in aqueous alkaline solution. Suitable solvents mentioned are methyl tert-butyl ether, toluene and xylene, the dielectric constant of these at 20°C being 2.6, 2.4 and, respectively, from 2.3 to 2.6.

The plastics described in DE 4234251 are colourless, rigid and slightly brittle and have a high refractive index n_D in the range from 1.602 to 1.608. The Abbe number is from 35 to 38. These plastics too, therefore, have only limited suitability for spectacle lenses. Again, this specification gives no information concerning the glass transition temperature of the plastics.

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The specification WO 03/011925 describes the polymerization of thiomethacrylates with polyethylene glycol derivatives. The resultant plastics may be used, inter alia, for producing optical lenses. A
5 disadvantage of these lenses is their mechanical properties. In particular, for example, impact strength is insufficient for many requirements.

In the light of the prior art, it was then an object of
10 the present invention to provide mixtures for preparing transparent plastics which are suitable as a material for optical lenses, where the plastics have ideal mechanical properties, in particular high impact strength, together with a high refractive index,
15 preferably greater than 1.59, and a maximum Abbe number, preferably greater than 36. In particular, it should be possible to produce plastics spectacle lenses which have a low level of dispersion and no colouring at the edges.

20 The present invention was also based on the object of providing a starting material composition for preparing a high-transparency plastic with improved mechanical properties, even at temperatures above room
25 temperature. In particular, the glass transition temperature of the inventive plastic should be maximized, preferably being above 80.0°C.

It was therefore an object of the present invention to
30 provide a high-transparency plastic which can be prepared from the starting material composition in a simple manner, on an industrial scale and at low cost. In particular, it should be obtainable via free-radical polymerization from a mixture which is flowable at
35 atmospheric pressure and temperatures in the range from 20.0 to 80.0°C.

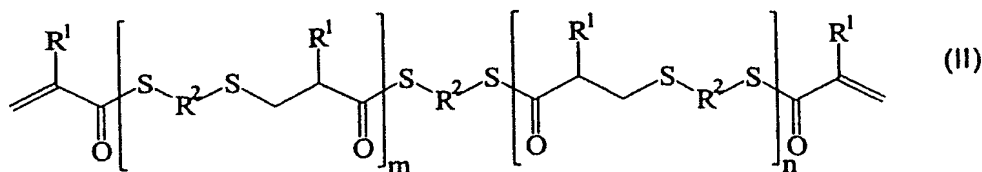
Another object on which the present invention was based was to provide application sectors and possible uses

for the inventive high-transparency plastic.

A mixture with all of the features of Patent Claim 1 achieves these objects, and also achieves other objects which, although not explicitly mentioned, are readily derivable or deducible from the circumstances discussed in the introduction. Advantageous modifications of the inventive mixture are protected in the subclaims dependent on Claim 1. Other subject matters claimed are the high-transparency plastics obtainable from the inventive mixtures, and also a process for their preparation. The use claim protects a preferred use of the inventive high-transparency plastic. Another product claim describes an optical, preferably ophthalmic, lens which comprises the inventive high-transparency plastic.

Mixtures encompassing

a) a prepolymer prepared from compounds of the formula (I) and (II)



where each R^1 , independently of the others, is hydrogen or a methyl radical,
each R^2 , independently of the others, is a linear or branched, aliphatic or cycloaliphatic radical, or a substituted or unsubstituted aromatic or heteroaromatic radical, and each of m and n , independently of the other, is a whole number greater than or equal to 0, where $m + n > 0$, and

from alkyl dithiols or from polythiols, preferably compounds of the formula (III)



where R^3 can be identical or different from the definition given for R^2

b) at least one monomer (A) capable of free-radical polymerization and having at least 2 methacrylate groups, and

c) aromatic vinyl compounds,

which are suitable for preparing transparent plastics, and which have excellent mechanical and optical properties. The mixtures may, if appropriate, comprise

d) a monomer capable of free-radical polymerization and having at least two terminal olefinic groups whose reactivity differs, for example a bifunctional monomer having a methacrylate end group and a vinyl end group, and/or

e) at least one ethylenically unsaturated monomer (B), preferably from the group of the methacrylates, particularly preferably 2-hydroxyethyl methacrylate.

The inventive transparent plastic has a previously unknown combination of exceptional properties, such as high refractive index, high Abbe number, good impact strength, and also high glass transition temperature. The corresponding plastics spectacle lenses have a low level of dispersion; no colouring at the edges is observed.

The inventive transparent plastic also has further advantages. Among these are:

⇒ Since the inventive plastic has high refractive

index, there is no requirement for reinforcement and therefore thickening of the centre and of the edges of corresponding plastics spectacle lenses, and there is a marked increase in the wearer comfort provided by these spectacles, due to the comparatively low weight.

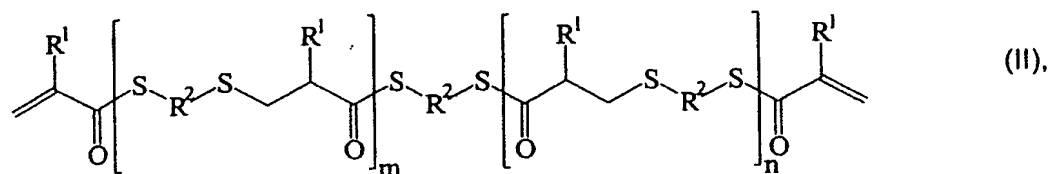
⇒ The very good impact strength of the inventive plastic protects the corresponding plastics spectacle lenses from the "risks of everyday life". In particular in the case of thin spectacle lenses, it is very unlikely that mechanical forces will cause impairment or irreparable damage.

⇒ The glass transition temperature of the inventive high-transparency plastic is high, preferably above 80.0°C, and up to this temperature the plastic therefore retains its exceptional mechanical properties, in particular high impact strength, and its hardness.

⇒ The inventive high-transparency plastic can be prepared via free-radical copolymerization in a simple manner, on an industrial scale, at low cost, of a monomer mixture which is preferably flowable at atmospheric pressure and temperatures in the range from 20.0 to 80.0°C.

⇒ The underlying monomer mixture can likewise be prepared in a simple manner, on an industrial scale, and at low cost.

The prepolymer of the present invention encompasses compounds of the formula (I) and/or (II) and (III)



5

where each R^1 , independently of the others, is hydrogen or a methyl radical, preferably a methyl radical, and each R^2 , independently of the others, is a linear or branched, aliphatic or cycloaliphatic radical, or a substituted or unsubstituted aromatic or heteroaromatic radical, where the radical R^2 may preferably encompass from 1 to 100, in particular from 1 to 20, carbon atoms and

each radical R^3 , irrespective of R^2 , is a linear or branched, aliphatic or cycloaliphatic radical or a substituted or unsubstituted aromatic or heteroaromatic radical, where the radical R^3 can preferably encompass from 1 to 100, in particular from 1 to 20, carbon atoms.

20

By way of example, among the preferred linear or branched, aliphatic or cycloaliphatic radicals are the methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene, or cyclohexylene group.

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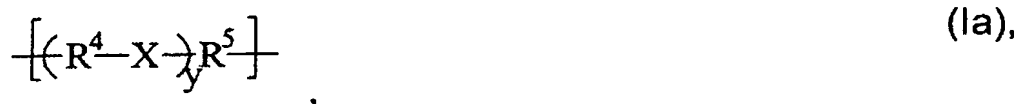
Among the preferred divalent aromatic or heteroaromatic radicals are in particular groups which derive from benzene, from naphthalene, from biphenyl, from diphenyl ether, from diphenylmethane, from diphenyldimethylmethane, from bisphenone, from diphenyl sulphone, from

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quinoline, from pyridine, from anthracene, and from phenanthrene. For the purposes of the present invention, cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic radicals.

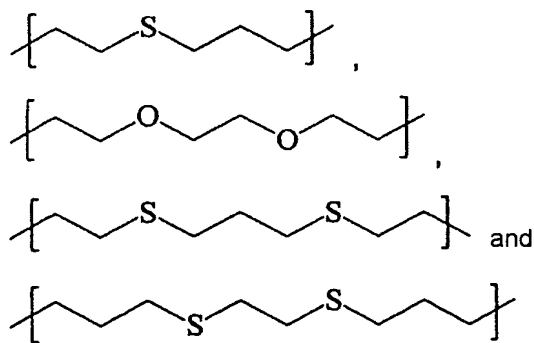
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The radical R^2 or R^3 also encompasses radicals of the formula



where each R^4 , independently of the others, is a linear or branched, aliphatic or cycloaliphatic radical, e.g. a methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group. Each radical X, independently of the others, is oxygen or sulphur, and the radical R^5 is a linear or branched, aliphatic or cycloaliphatic radical, e.g. a methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group. For the purposes of the present invention, cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic radicals. Y is a whole number from 1 to 10, in particular 1, 2, 3 or 4.

Preferred radicals of the formula (Ia) encompass:



25

The radical R^2 is preferably an aliphatic radical having from 1 to 10 carbon atoms, preferably a linear

aliphatic radical having from 2 to 8 carbon atoms.
Each of the indices m and n, independently of the others, is a whole number greater than or equal to 0, such as 0, 1, 2, 3, 4, 5 or 6. The sum $m + n$ here is
5 greater than 0, preferably in the range from 1 to 6, advantageously in the range from 1 to 4, in particular 1, 2 or 3.

To prepare the prepolymer, each of the compounds of the
10 formula (I) and (II), and also each of the compounds of the formula (III), may be used individually or else in the form of a mixture of two or more compounds of the formula (I), (II) and, respectively, (III).

15 The relative proportions of the compounds of the formula (I), (II) and (III) in the inventive monomer mixture may in principle be as desired, and they can be utilized to "tailor" the property profile of the
20 inventive plastic to the demands of the application. By way of example, it can be extremely advantageous for the monomer mixture to comprise a marked excess of compound(s) of the formula (I) or compound(s) of the formula (II) or compound(s) of the formula (III), based in each case on the total amount of compounds of the
25 formula (I), (II) and (III) in the prepolymer.

However, for the purposes of the present invention it is particularly advantageous for the mixture to comprise more than 10 mol%, preferably more than
30 12 mol%, in particular more than 14 mol%, based on the total amount of the compounds of the formula (I) and (II), of compounds of the formula (II) where $m + n = 2$. If R^2 is an ethylene radical, the proportion by weight of (II) where $m + n = 2$ in the mixture is more than
35 10%, in particular more than 15%.

It is moreover particularly advantageous according to the invention to use mixtures which comprise more than 5.8 mol%, advantageously more than 6.5 mol%, in

particular more than 7.5 mol%, based on the total amount of the compounds of the formula (I) and (II), of compounds of the formula (II) where $m + n = 3$. This corresponds to a proportion by weight of (II) of at least 6% if R^2 is an ethylene radical, where $m + n = 3$.

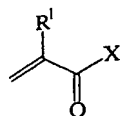
The proportion of the compounds (I) is preferably from 0.1 to 50.0 mol%, advantageously from 10.0 to 45.0 mol%, in particular from 20.0 to 35.0 mol%, based on the total amount of the compounds of the formula (I) and (II), corresponding to a preferred range for the preparation by weight of the compound (I) of from 15 to 40% if R^2 is an ethylene radical. The proportion of the compounds (II) where $m + n = 1$ is preferably from 1 to 40.0 mol%, advantageously from 5 to 35.0 mol%, in particular from 10 to 30 mol%, based on the total amount of the compounds of the formula (I) and (II). This corresponds to a preferred proportion by weight of the compounds (II) where $m + n = 1$ of from 10 to 45% if R^2 is an ethylene radical. The proportion of the compounds (II) where $m + n > 3$ is preferably greater than 0 mol%, advantageously greater than 1 mol%, in particular greater than 2 mol%, based on the total amount of the compounds of the formula (I) and (II). If R^2 is an ethylene radical, the proportion by weight for compounds (II) where $m + n > 3$ in the mixture is more than 2%, in particular more than 5%.

The proportion of the compounds (III) in the prepolymer is preferably from 1 to 55.0 mol%, in particular from 10.0 to 50.0 mol%, based on the total amount of the compounds of the formula (I), (II) and (III). If, in a specific case, R^3 is a dimercaptodioxaoctane radical, then the proportion by weight of (III) in the prepolymer, based on the total amount of the compounds (I), (II) and (III) is more than 0.5%, preferably more than 5%.

Processes for preparing the compounds of the formula

(I) and (II) are known to the person skilled in the art, for example from DE 4234251, the disclosure of which is expressly incorporated herein by way of reference. However, for the purposes of the present invention, it has proven very particularly advantageous to prepare a mixture of the compounds of the formula (I) and (II) via a process in which from 1.0 to < 2.0 mol, preferably from 1.1 to 1.8 mol, advantageously from 1.2 to 1.6 mol, in particular from 1.2 to 1.5 mol, of at least one compound of the formula (IV)

(IV)

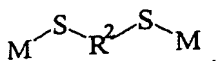


are reacted with one mole of at least one polythiol of the formula (V)

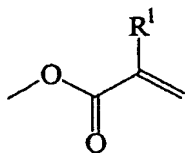
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(V)

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The radical X is halogen, in particular chlorine or bromine, or a radical



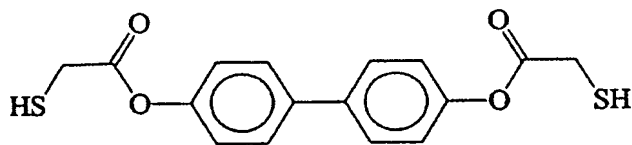
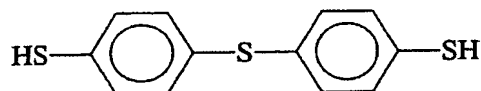
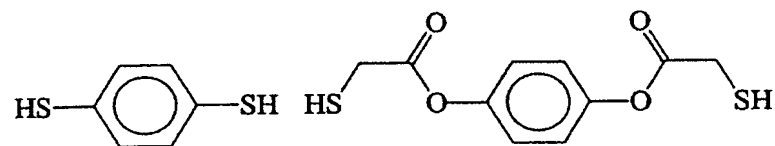
this means that the compounds of the formula (IV) encompass, inter alia, acryloyl chloride, methacryloyl chloride, acrylic anhydride and methacrylic anhydride, particular preference being given to the use of acrylic anhydride, methacrylic anhydride or mixtures of the two.

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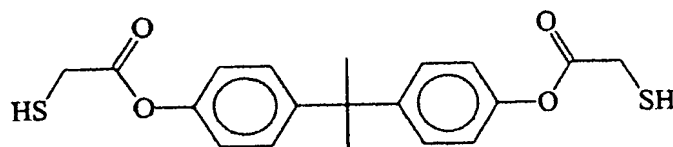
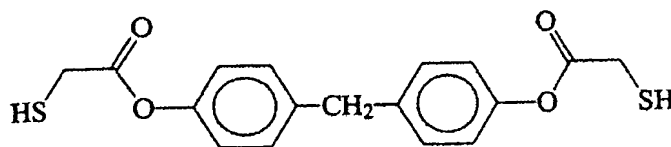
Each M indicates, independently of the other, hydrogen or a metal cation. Preferred metal cations derive from elements whose electronegativity is smaller than 2.0,

advantageously smaller than 1.5, particular preference being given to alkali metal cations, in particular Na^+ , K^+ , Rb^+ , Cs^+ and alkaline earth metal cations, in particular Mg^{2+} , Ca^{2+} , Sr^{2+} , Ba^{2+} . Very particularly
5 advantageous results may be achieved using the metal cations Na^+ and K^+ .

Polythiols of the formula (V) particularly suitable in this context encompass 1,2-ethanedithiol, 1,2-propanedithiol, 10 dithiol, 1,3-propanedithiol, 1,2-butanedithiol, 1,3-butanedithiol, 1,4-butanedithiol, 2-methylpropane-1,2-dithiol, 2-methylpropane-1,3-dithiol, 3,6-dioxo-1,8-octanedithiol (dimercaptodioxaoctane = DMDO), ethylcyclohexyl dimercaptans obtainable via reaction of
15 4-ethenylcyclohexene with hydrogen sulphite, ortho-bis(mercaptomethyl)benzene, meta-bis(mercaptomethyl)benzene, para-bis(mercaptomethyl)-benzene, compounds of the formula (V)



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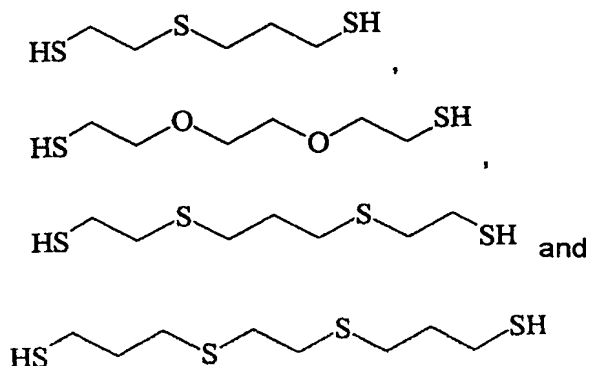


and also compounds of the formula



5 where each R^4 , independently of the others, is a linear or branched, aliphatic or cycloaliphatic radical, such as a methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group. For the purposes of the present invention, 10 cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic radicals. Each radical X, independently of the others, is oxygen or sulphur, and the radical R^5 is a linear or branched, aliphatic or cycloaliphatic radical, such as a methylene, ethylene, 15 propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group. For the purposes of the present invention, cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic radicals. y is a whole number from 1 to 10, in 20 particular 1, 2, 3 or 4.

Preferred compounds of the formula (Va) encompass:



25

For the purposes of one particularly preferred embodiment of this process, the compound used of the formula (V) comprises 1,2-ethanedithiol.

30

This process reacts the compound(s) of the formula (IV) in at least one inert, organic solvent S, and the compound(s) of the formula (V) in aqueous alkaline solution, the expression "inert, organic solvent" applying to organic solvents which do not react with the compounds present in the reaction system under the respective reaction conditions.

It is preferable for at least one solvent S to have a relative dielectric constant > 2.6 , preferably > 3.0 , advantageously > 4.0 , in particular > 5.0 , in each case measured at 20°C . In this context, the relative dielectric constant indicates a dimensionless value stating the factor by which the capacitance C of a (theoretical) capacitor located within a vacuum increases when substances with dielectric properties, known as dielectrics, are introduced between the plates. This value is measured at 20°C and extrapolated to low frequencies ($\omega \rightarrow 0$). For further details reference is made to the familiar technical literature, in particular to Ullmann Encyklopädie der technischen Chemie, [Ullmann's Encyclopaedia of Industrial Chemistry] Volume 2/1 Anwendung physikalischer und physikalisch-chemischer Methoden im Laboratorium [Application of physical and physico-chemical methods in the laboratory], headword: Dielektrizitätskonstante [Dielectric constant], pp. 455-479. Dielectric values of solvents are given, inter alia, in Handbook of Chemistry and Physics, 71st edition, CRC Press, Boca Raton, Ann Arbor, Boston, 1990-1991, pp. 8-44, 8-46 and 9-9 to 9-12.

For the purposes of this process it is moreover particularly advantageous for the solvent and the aqueous solution to form two phases during the reaction and not to be capable of homogeneous mixing. To this end, the water solubility value for the solvent, measured at 20°C , is preferably smaller than 10 g of water, based on 100 g of solvent.

Solvents S preferred according to the invention encompass

aliphatic ethers, such as diethyl ether (4.335), dipropyl ether, diisopropyl ether;

5 cycloaliphatic ethers, such as tetrahydrofuran (7.6);

aliphatic esters, such as methyl formate (8.5), ethyl formate, propyl formate, methyl acetate, ethyl acetate, n-butyl acetate (5.01), methyl propionate, methyl butyrate (5.6), ethyl butyrate, 2-methoxyethyl acetate;

10 aromatic esters, such as benzyl acetate, dimethyl phthalate, methyl benzoate (6.59), ethyl benzoate (6.02), methyl salicylate, ethyl salicylate, phenyl acetate (5.23);

aliphatic ketones, such as acetone, methyl ethyl ketone
15 (18.5), 2-pentanone (15.4), 3-pentanone (17.0), methyl isoamyl ketone, methyl isobutyl ketone (13.1);

aromatic ketones, such as acetophenone;

nitroaromatics, such as nitrobenzene, o-nitrotoluene (27.4), m-nitrotoluene (23), p-nitrotoluene;

20 halogenated aromatics, such as chlorobenzene (5.708), o-chlorotoluene (4.45), m-chlorotoluene (5.55), p-chlorotoluene (6.08), o-dichlorobenzene, m-dichlorobenzene;

heteroaromatics, such as pyridine, 2-methylpyridine
25 (9.8), quinoline, isoquinoline; and mixtures of these compounds, the data in brackets being the respective associated relative dielectric constants at 20°C.

Compounds very particularly suitable here for the
30 purposes of the present invention are aliphatic esters and cycloaliphatic ethers, in particular ethyl acetate and tetrahydrofuran.

For the purposes of the present process, it is possible
35 either to use the solvent S alone or else to use a

solvent mixture, in which case it is not necessary that all of the solvents present in the mixture comply with the abovementioned dielectric criterion. By way of example, according to the invention it is also possible
5 to use tetrahydrofuran/cyclohexane mixtures. However, it has proven advantageous for the solvent mixture to have a relative dielectric constant > 2.6 , preferably > 3.0 , advantageously > 4.0 , in particular > 5.0 , in each case measured at 20°C . Particularly advantageous
10 results can be achieved using solvent mixtures which comprise only solvents whose relative dielectric constant is > 2.6 , preferably > 3.0 , advantageously > 4.0 , in particular > 5.0 , in each case measured at 20°C .

15 The aqueous alkaline solution of the compound(s) of the formula (V) preferably comprises from 1.1 to 1.5 val (equivalents) of at least one Bronsted base, based on the total amount of compound(s) of the formula (IV).
20 Preferred Bronsted bases for the purposes of the present invention encompass alkali metal hydroxides and alkaline earth metal hydroxides, in particular sodium hydroxide and potassium hydroxide.

25 In principle, any conceivable method may be used for the conduct of the reaction. By way of example, the compound(s) of the formula (IV) may form an initial charge in the solvent (mixture) S, and the aqueous alkaline solution of the compound(s) of the formula (V)
30 may be added stepwise or continuously. However, for the purposes of the present invention it has proven very particularly advantageous to meter the compound(s) of the formula (IV) in at least one inert, organic solvent S and the compound(s) of the formula (V) in
35 aqueous alkaline solution to the reaction vessel in parallel.

The reaction temperature may be varied widely, but the temperature is often in the range from 20.0 to 120.0°C ,

preferably in the range from 20.0 to 80.0°C. Similar considerations apply for the pressure at which the reaction is completed. The reaction may therefore take place either at subatmospheric pressure or else at
5 superatmospheric pressure. However, it is preferably carried out at atmospheric pressure. Although the reaction can also take place in air, it has proven very particularly advantageous for the purposes of the present invention to carry out the reaction under an
10 inert gas, preferably nitrogen and/or argon, preferably with a very small proportion of oxygen present.

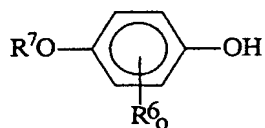
The reaction mixture is advantageously reacted in a further step with a Bronsted acid, preferably until the
15 pH of the aqueous solution at 20°C is below 7.0, advantageously below 6.0, in particular below 5.0. Acids which may be used in this context encompass inorganic mineral acids, such as hydrochloric acid, sulphuric acid, phosphoric acid, organic acids, such as
20 acetic acid, propionic acid, and acidic ion exchangers, in particular acidic synthetic resin ion exchangers, e.g. ®Dowex M-31 (H). The method which has proven very particularly successful here is the use of acidic synthetic resin ion exchangers loaded with at least
25 1.0 meq, preferably at least 2.0 meq, in particular at least 4.0 meq, of H⁺ ions, based on 1 g of dry ion exchanger, grain sizes of from 10 to 50 mesh and porosities in the range from 10 to 50%, based on the total volume of the ion exchanger.

30 In an advantageous method for isolating the compounds of the formula (I) and (II) the organic phase composed of the solvent S is separated off and, where appropriate, washed, and dried, and the solvent is
35 evaporated.

During the reaction of the compound(s) of the formula (IV) with the compound(s) of the formula (V) it is possible to add inhibitors which inhibit free-

radical polymerization of the (meth)acrylic groups during the reaction. These inhibitors are well-known to persons skilled in the art.

- 5 Use is mainly made of 1,4-dihydroxybenzenes. However, it is also possible to use dihydroxybenzenes having other substitution. These inhibitors can generally be represented by the general formula (VI)



(VI),

10

where

- 15 R⁶ is a linear or branched alkyl radical having from one to eight carbon atoms, halogen or aryl, preferably an alkyl radical having from one to four carbon atoms, particularly preferably methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, Cl, F or Br;

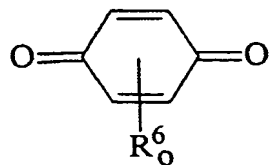
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o is a whole number in the range from one to four, preferably one or two; and

- 25 R⁷ is hydrogen, a linear or branched alkyl radical having from one to eight carbon atoms, or aryl, preferably an alkyl radical having from one to four carbon atoms, particularly preferably methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl or tert-butyl.

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However, it is also possible to use compounds whose parent compound is 1,4-benzoquinone. These may be described by the formula (VII)

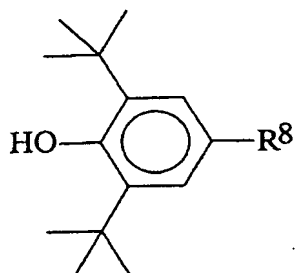


(VII)

where

5 R⁶ and o are as defined above.

Use is also made of phenols of the general structure (VIII)



(VIII)

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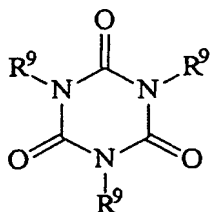
where

15 R⁸ is a linear or branched alkyl radical having from one to eight carbon atoms, aryl or aralkyl, propionic esters with mono- to tetrahydric alcohols, which may also contain heteroatoms, such as S, O and N, preferably an alkyl radical having from 1 to 4 carbon atoms, particularly preferably methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl.

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Another advantageous class of substance is represented by hindered phenols based on triazine derivatives of the formula (IX)

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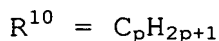


(IX)

where R^9 = a grouping of the formula (X)



5 where



where $p = 1$ or 2 .

10

Compounds used with particular success are 1,4-dihydroxybenzene, 4-methoxyphenol, 2,5-dichloro-3,6-dihydroxy-1,4-benzoquinone, 1,3,5-trimethyl-2,4,6-tris(3,5-di-tert-butyl-4-hydroxybenzyl)benzene, 2,6-di-
15 tert-butyl-4-methylphenol, 2,4-dimethyl-6-tert-butylphenol, 2,2-bis[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl-1-oxopropoxymethyl]1,3-propanediyl ester, 2,2'-thiodiethyl bis[3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionate], octadecyl 3-(3,5-di-tert-
20 butyl-4-hydroxyphenyl)propionate, 3,5-bis(1,1-dimethylethyl-2,2'-methylenebis(4-methyl-6-tert-butyl)phenol, tris(4-tert-butyl-3-hydroxy-2,6-dimethylbenzyl)-s-triazine-2,4,6-(1H,3H,5H)trione, tris(3,5-di-tert-butyl-4-hydroxy)-s-triazine-2,4,6-
25 (1H,3H,5H)trione or tert-butyl-3,5-dihydroxybenzene.

Based on the weight of the entire reaction mixture, the proportion of the inhibitors individually or in the form of a mixture is generally from 0.01 to 0.50% (by
30 weight), the concentration of the inhibitors preferably being selected in such a way as not to impair the DIN 55945 colour number. Many of these inhibitors are commercially available.

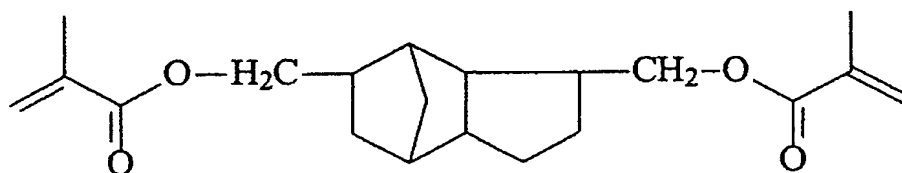
35 For the purposes of the present invention, the mixture

also comprises, alongside the prepolymer, prepared from compounds of the formula (I), (II) and (III), at least one monomer (A) capable of free-radical polymerization and having at least two terminal methacrylate groups.

5 Examples of these di(meth)acrylates are polyoxymethylene-(meth)acrylic acid derivatives and polyoxypropylene-(meth)acrylic acid derivatives, e.g. triethylene glycol (meth)acrylate, tetraethylene glycol
10 (meth)acrylate, tetrapropylene glycol (meth)acrylate, and also 1,4-butanediol di(meth)acrylate, diethylene glycol di(meth)acrylate, dipropylene glycol di(meth)acrylate, triethylene glycol di(meth)acrylate, tripropylene glycol di(meth)acrylate. Tetraethylene
15 glycol di(meth)acrylate, tetrapropylene glycol di(meth)acrylate, polyethylene glycol di(meth)acrylate (preferably having weight-average molar masses in the range from 200 to 5 000 000 g/mol, advantageously in the range from 200 to 25 000 g/mol, in particular in
20 the range from 200 to 1000 g/mol), polypropylene glycol di(meth)acrylate (preferably with weight-average molar masses in the range from 200 to 5 000 000 g/mol, advantageously in the range from 250 to 4000 g/mol, in particular in the range from 250 to 1000 g/mol), 2,2'-
25 thiodiethanol di(meth)acrylate (thiodiglycol di(meth)acrylate),

3,9-di(meth)acryloyloxymethyltricyclo[5.2.1.0(2.6)]-
decane, in particular

30



3,8-di(meth)acryloyloymethyltricyclo[5.2.1.0(2.6)]-
decane

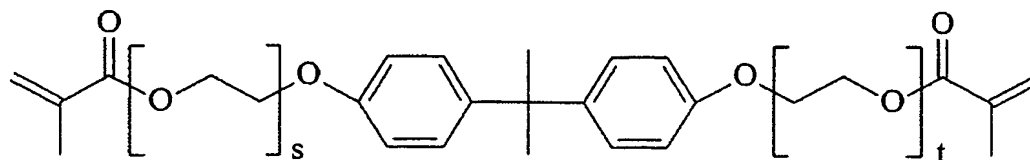
35 4,8-di(meth)acryloyloymethyltricyclo[5.2.1.0(2.6)]-

decane,

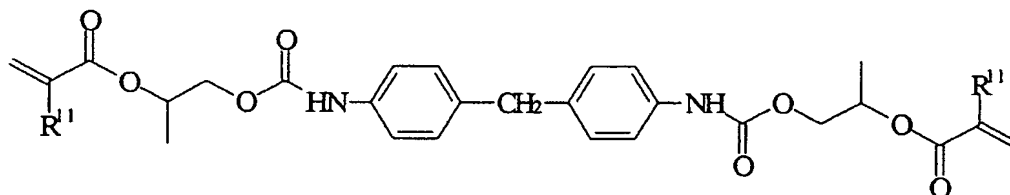
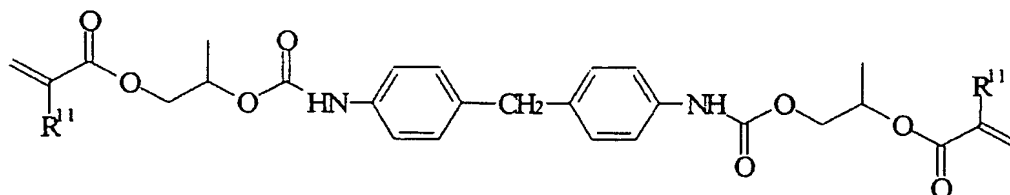
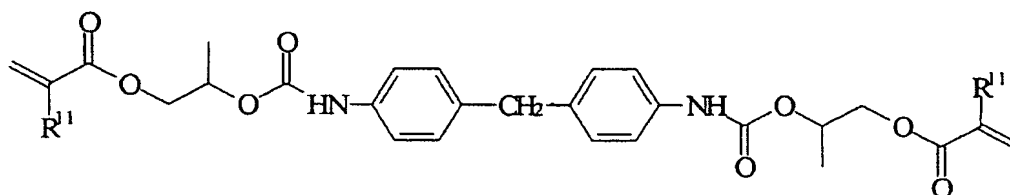
4,9-di(meth)acryloyloymethyltricyclo[5.2.1.0(2.6)]-

decane, ethoxylated bisphenol A di(meth)acrylate, in particular

5

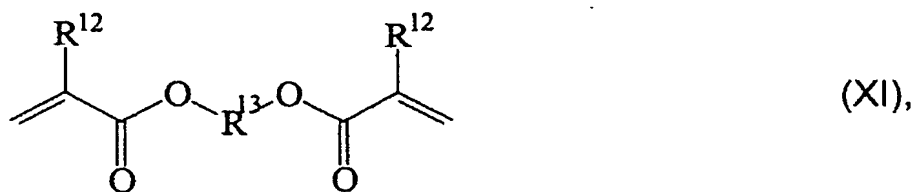


where s and t are greater than or equal to zero, and the sum of s and t is preferably in the range from 1 to 10 30, in particular in the range from 2 to 10, and di(meth)acrylates obtainable via reaction of diisocyanates with 2 equivalents of hydroxyalkyl (meth)acrylate, in particular



15 where each radical R^{11} , independently of the others, is hydrogen or a methyl radical, tri(meth)acrylates, such as trimethylolpropane tri(meth)acrylate and glycerol tri(meth)acrylate, or else (meth)acrylates of ethoxylated or propoxylated 20 glycerol, of trimethylolpropane, or of other alcohols having more than 2 hydroxy groups.

Di(meth)acrylates of the formula (XI) have proven particularly successful as monomer (A).



5 Each R^{12} here, independently of the others, is hydrogen or methyl. R^{13} indicates a linear or branched alkyl or cycloalkyl radical, or an aromatic radical preferably having from 1 to 100, with preference from 1 to 40,
10 preferably from 1 to 20, advantageously from 1 to 8, in particular from 1 to 6, carbon atoms, e.g. a methyl, ethyl, propyl, isopropyl, n-butyl, isobutyl, tert-butyl, cyclopentyl, cyclohexyl, or phenyl group. For the purposes of the present invention, cycloaliphatic
15 radicals here also encompass bi-, tri-, and polycyclic aliphatic radicals. Linear or branched alkyl or cycloalkyl radicals having from 1 to 6 carbon atoms are very particularly preferred as R^{18} .

20 The radical R^{13} is preferably a linear or branched, aliphatic or cycloaliphatic radical, such as a methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group, or a radical of the general formula

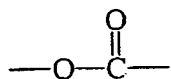
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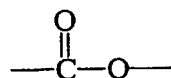
where the radical R^{15} is a linear or branched, aliphatic or cycloaliphatic radical, or a substituted or
30 unsubstituted aromatic or heteroaromatic radical, e.g. a methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group, or divalent aromatic or heteroaromatic groups

which derive from benzene, from naphthalene, from biphenyl, from diphenyl ether, from diphenylmethane, from diphenyldimethylmethane, from bisphenone, from diphenyl sulphone, from quinoline, from pyridine, from anthracene, or from phenanthrene. For the purposes of the present invention, cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic radicals. Each radical R^{14} , independently of the others, is a linear or branched, aliphatic or cycloaliphatic radical or a substituted or unsubstituted aromatic or heteroaromatic radical, e.g. a methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group, or divalent aromatic or heteroaromatic groups which derive from benzene, from naphthalene, from biphenyl, from diphenyl ether, from diphenylmethane, from diphenyldimethylmethane, from bisphenone, from diphenyl sulphone, from quinoline, from pyridine, from anthracene, or from phenanthrene. For the purposes of the present invention, cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic radicals. Each radical X^I , independently of the others, is oxygen, sulphur, an ester group of the general formula (XIb), (XIc),

25



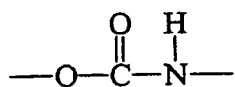
(XIb)



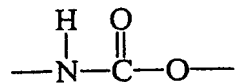
(XIc)

a urethane group of the general formula (XIId), (XIE), (XIIf) or (XIg),

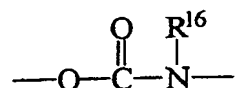
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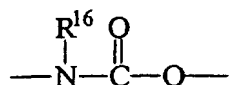
(XId)



(XIe)

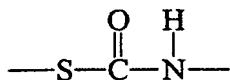


(XI f)

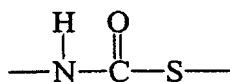


(XIg)

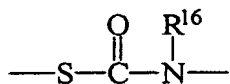
a thiourethane group of the general formula (XIh),
5 (XIi), (XIj) or (XI k),



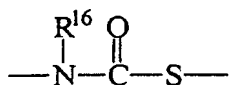
(XIh)



(XIi)

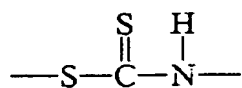


(XIj)

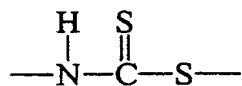


(XV k)

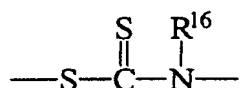
a dithiourethane group of the general formula (XI l),
10 (XI m), (XI n) or (XI o)



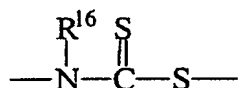
(XII)



(XIm)



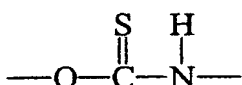
(XIn)



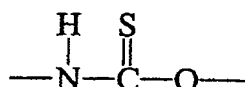
(XIo)

or a thiocarbamate group of the general formula (XIp), (XIq), (XIr) or (XIs)

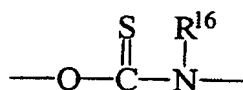
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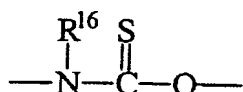
(XIp)



(XIq)



(XIr)

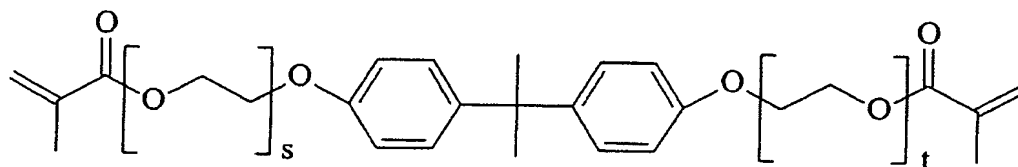


(XIs)

- 10 preferably oxygen, where the radical R^{16} is a linear or branched, aliphatic or cycloaliphatic radical or a substituted or unsubstituted aromatic or heteroaromatic radical, e.g. a methyl, ethyl, propyl, isopropyl, n-butyl, isobutyl, tert-butyl or cyclohexyl group, or
- 15 monovalent aromatic or heteroaromatic groups derived from benzene, from naphthalene, from biphenyl, from diphenyl ether, from diphenylmethane, from

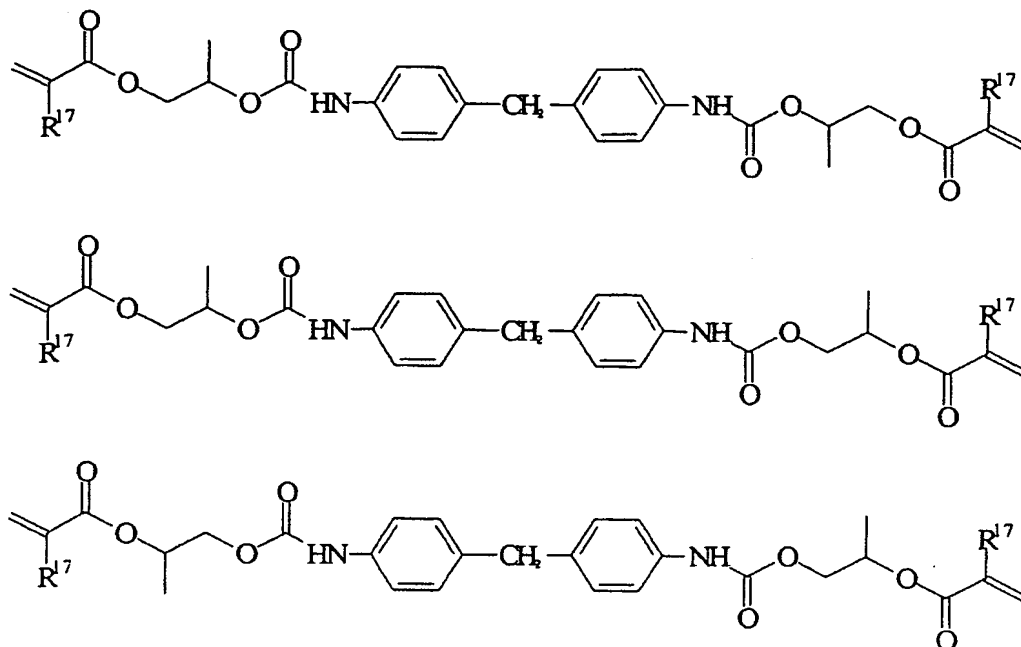
diphenyldimethylmethane, from bisphenone, from diphenyl sulphone, from quinoline, from pyridine, from anthracene, or from phenanthrene. For the purposes of the present invention, cycloaliphatic radicals here
5 also encompass bi-, tri- and polycyclic aliphatic radicals. z is a whole number from 1 to 1000, advantageously from 1 to 100, in particular from 1 to 25.

10 Particularly preferred di(meth)acrylates of the formula (XI) encompass ethylene glycol di(meth)acrylate, ethoxylated bisphenol A di(meth)acrylate, in particular



15

where s and t are greater than or equal to zero and the sum of s and t is preferably in the range from 1 to 20, in particular in the range from 2 to 10, and di(meth)acrylates obtainable via reaction of
20 diisocyanates with 2 equivalents of hydroxyalkyl (meth)acrylate, in particular



where each radical R^{17} , independently of the others, is hydrogen or a methyl radical,

5 3,8-di(meth)acryloyloxymethyltricyclo[5.2.1.0(2.6)]-decane,

3,9-di(meth)acryloyloxymethyltricyclo[5.2.1.0(2.6)]-decane,

10 4,8-di(meth)acryloyloxymethyltricyclo[5.2.1.0(2.6)]-decane,

4,9-di(meth)acryloyloxymethyltricyclo[5.2.1.0(2.6)]-decane,

thiodiglycol di(meth)acrylate, polypropylene glycol di(meth)acrylate, preferably with a weight-average molar mass in the range from 200 to 1000 g/mol, and/or polyethylene glycol di(meth)acrylate, preferably with a weight-average molar mass in the range from 200 to 1000 g/mol. Particular preference is given here to the dimethacrylates of the compounds mentioned. Very particularly advantageous results are achieved using polyethylene glycol dimethacrylate, preferably with a weight-average molar mass in the range from 200 to 1000 g/mol.

25 The proportion of monomer (A) is from 2 to 50% by

weight, in particular from 10 to 30% by weight, based on all of the monomers used in the mixture.

For the purposes of the present invention, the mixture
5 also comprises, alongside the prepolymer composed of the compounds of the formula (I), (II), and (III), and at least one monomer (A) capable of free-radical polymerization, an aromatic vinyl compound.

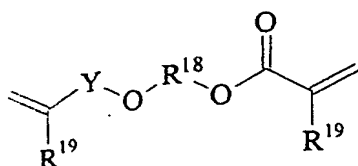
10 Among the aromatic vinyl compounds, preference is given to the use of styrenes, substituted styrenes having an alkyl substituent in the side chain, e.g. α -methylstyrene and α -ethylstyrene, substituted styrenes having an alkyl substituent on the ring, e.g.
15 vinyltoluene and p-methylstyrene, halogenated styrenes, such as monochlorostyrenes, dichlorostyrenes, tribromostyrenes, and tetrabromostyrenes,

and also to dienes, such as 1,2-divinylbenzene,
20 1,3-divinylbenzene, 1,4-divinylbenzene, 1,2-diisopropenylbenzene, 1,3-diisopropenylbenzene, and 1,4-diisopropenylbenzene.

The proportion of the aromatic vinyl compounds is from
25 5 to 40% by weight, preferably from 10 to 30% by weight, particularly preferably from 15 to 25% by weight, based on the total amount of the compounds of the formula (I), (II), and (III) used in the prepolymer, the monomer (A) capable of free-radical
30 polymerization, and the aromatic vinyl compounds and other monomers optionally used.

Surprisingly, the addition of monomer (A) and the aromatic vinyl compound improves the mechanical
35 properties of the inventive plastics material without adversely affecting its optical properties. In many instances, a favourable effect on optical properties is found.

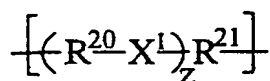
According to one particular aspect of the present invention, compounds, preferably molecules having a linear structure and varying chain lengths (asymmetric crosslinking agents) of the general formula (XII) may
5 be present



(XII),

where the radical R^{19} is independently a hydrogen atom,
10 a fluorine atom and/or a methyl group, the radical R^{18} is a connecting group, preferably encompassing from 1 to 1000, in particular from 2 to 100, carbon atoms and the radical Y is a bond or a connecting group having
15 from 0 to 1000 carbon atoms, in particular from 1 to 1000 carbon atoms, and preferably from 1 to 100 carbon atoms. The length of the molecule can be varied by way of the molecular component R^{18} . Compounds of the formula (XII) have, at one end of the molecule, a terminal (meth)acrylate function, and at the other end
20 have a terminal group other than a methacrylate function. Among the preferred groups Y are in particular a bond (vinyl group), a CH_2 group (allyl group), and also aliphatic or aromatic groups having from 1 to 20 carbon atoms, for example a benzene-
25 derived group, the aliphatic or aromatic groups particularly preferably containing a urethane group.

The radical R^{18} is preferably a linear or branched, aliphatic or cycloaliphatic radical, such as a
30 methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group, or a radical of the general formula



(XIIa),

where the radical R^{21} is a linear or branched, aliphatic or cycloaliphatic radical or a substituted or unsubstituted aromatic or heteroaromatic radical, e.g.

5 a methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group, or divalent aromatic or heteroaromatic groups which derive from benzene, from naphthalene, from biphenyl, from diphenyl ether, from diphenylmethane,

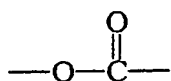
10 from diphenyldimethylmethane, from bisphenone, from diphenyl sulphone, from quinoline, from pyridine, from anthracene or from phenanthrene. For the purposes of the present invention, cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic

15 radicals. Each radical R^{20} here, independently of the others, is a linear or branched, aliphatic or cycloaliphatic radical or a substituted or unsubstituted aromatic or heteroaromatic radical, e.g.

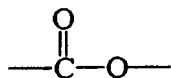
20 a methylene, ethylene, propylene, isopropylene, n-butylene, isobutylene, tert-butylene or cyclohexylene group, or divalent aromatic or heteroaromatic groups which derive from benzene, from naphthalene, from biphenyl, from diphenyl ether, from diphenylmethane, from diphenyldimethylmethane, from bisphenone, from

25 diphenyl sulphone, from quinoline, from pyridine, from anthracene or from phenanthrene. For the purposes of the present invention, cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic radicals. Each radical X^1 , independently of the others,

30 is oxygen, sulphur, an ester group of the general formula (XIIf), (XIIf),

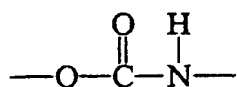


(XIIf)

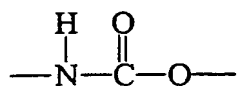


(XIIf)

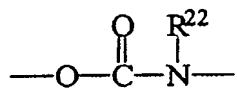
a urethane group of the general formula (XIId), (XIIE), (XIIf) or (XIIG),



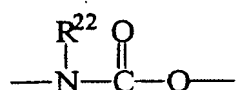
(XIId)



(XIIE)



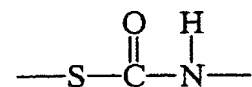
(XIIf)



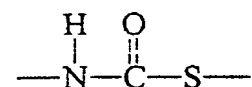
(XIIG)

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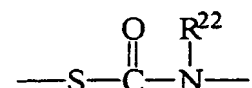
a thiourethane group of the general formula (XIIH), (XIIi), (XIIj) or (XIIk),



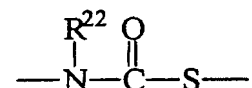
(XIIH)



(XIIi)

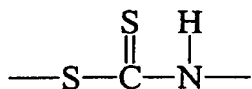


(XIIj)

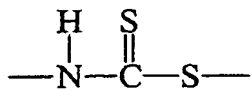


(XIIk)

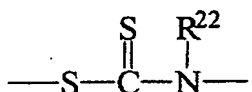
10 a dithiourethane group of the general formula (XIIl), (XIIIm), (XIIn) or (XIIo)



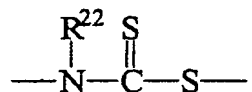
(XIII)



(XIIm)

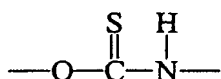


(XIIn)

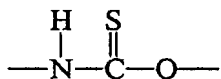


(XIlo)

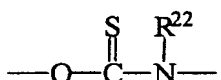
or a thiocarbamate group of the general formula (XIIp),
 5 (XIIq), (XIIr) or (XIIs)



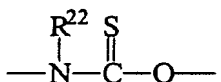
(XIIp)



(XIIq)



(XIIr)

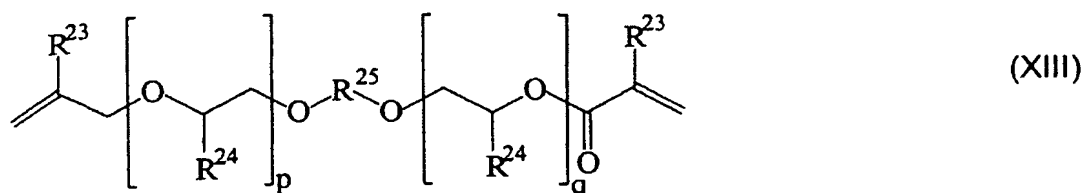


(XIIs)

preferably oxygen, where the radical R^{22} is a linear or
 10 branched, aliphatic or cycloaliphatic radical or a
 substituted or unsubstituted aromatic or heteroaromatic
 radical, e.g. a methyl, ethyl, propyl, isopropyl,
 n-butyl, isobutyl, tert-butyl or cyclohexyl group, or
 monovalent aromatic or heteroaromatic groups derived
 15 from benzene, from naphthalene, from biphenyl, from
 diphenyl ether, from diphenylmethane, from
 diphenyldimethylmethane, from bisphenone, from diphenyl
 sulphone, from quinoline, from pyridine, from

anthracene, or from phenanthrene. For the purposes of the present invention, cycloaliphatic radicals here also encompass bi-, tri- and polycyclic aliphatic radicals. z is a whole number from 1 to 1000, advantageously from 1 to 100, in particular from 1 to 25.

In one particular embodiment of the formula (XII) the compounds comprise those of the formula (XIII)



and/or of the formula (XIV),

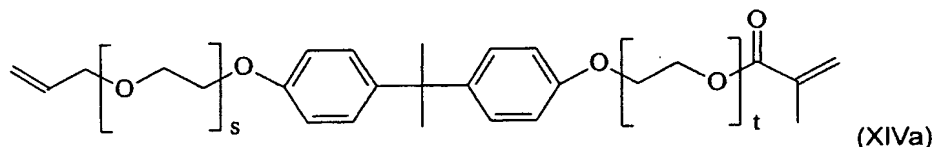


where each of the radicals R^{23} and R^{24} , independently of the other, is hydrogen or a methyl radical, and the radical R^{25} is a linear or branched, aliphatic or cycloaliphatic divalent radical, or a substituted or unsubstituted aromatic or heteroaromatic divalent radical. Preferred radicals have been described above.

The length of the chain may be influenced via variation of the number of polyalkylene oxide units, preferably polyethylene glycol units. Compounds of the formula (XIII) and (XIV) which have proven particularly suitable for the method described here of achieving the object have numbers of polyoxyalkylene oxide units r, p and q which are, independently of the others, from 1 to 40, preferably from 5 to 20, in particular from 7 to 15

and particularly preferably from 8 to 12.

Asymmetric crosslinking agents which are very particularly preferred according to the invention encompass compounds of the formula (XIV), in particular



where s and t are greater than or equal to zero and the sum s + t is preferably in the range from 1 to 20, in particular in the range from 2 to 10, and compounds of the formula (XIII), in particular



where s and t are greater than or equal to zero and the sum s + t is preferably in the range from 1 to 20, in particular in the range from 2 to 10.

According to one particular aspect, the mixture preferably comprises from 0.5 to 40% by weight, in particular from 5 to 15% by weight, of compounds of the formula (XII) and/or (XIII), based on the total weight of the monomer mixture.

For the purposes of one particularly preferred embodiment of the present invention, the inventive mixture also comprises at least one ethylenically unsaturated monomer (B). These monomers (B) differ from the asymmetric compounds of the formulae (XIII) and (XIV), and from the monomers (A) and the thio(meth)acrylates of the formulae (I) and/or (II). The monomers (B) are known to persons skilled in the

art and are preferably copolymerizable with the monomers (A) and the thio(meth)acrylates of the formulae (I) and/or (II). Among these monomers (B) are in particular

5

nitriles of (meth)acrylic acid and other nitrogen-containing methacrylates, such as methacryloylamidoacetonitrile, 2-methacryloyloxyethylmethylcyanamide, cyanomethyl methacrylate;

10

(meth)acrylates which derive from saturated alcohols, e.g. methyl (meth)acrylate, ethyl (meth)acrylate, n-propyl (meth)acrylate, isopropyl (meth)acrylate, n-butyl (meth)acrylate, sec-butyl (meth)acrylate, tert-butyl (meth)acrylate, pentyl (meth)acrylate, hexyl

15

(meth)acrylate, heptyl (meth)acrylate, 2-ethylhexyl (meth)acrylate, octyl (meth)acrylate, nonyl (meth)acrylate, isooctyl (meth)acrylate, isononyl (meth)acrylate, 2-tert-butylheptyl (meth)acrylate, 3-isopropylheptyl (meth)acrylate, decyl (meth)acrylate, undecyl (meth)acrylate, 5-methylundecyl (meth)acrylate, dodecyl (meth)acrylate, 2-methyldodecyl (meth)acrylate, tridecyl (meth)acrylate, 5-methyltridecyl

20

(meth)acrylate, tetradecyl (meth)acrylate, pentadecyl (meth)acrylate, hexadecyl (meth)acrylate, 2-methylhexadecyl (meth)acrylate, heptadecyl (meth)acrylate, 5-isopropylheptadecyl (meth)acrylate, 4-tert-butyloctadecyl (meth)acrylate, 5-ethyloctadecyl (meth)acrylate, 3-isopropyloctadecyl (meth)acrylate, octadecyl (meth)acrylate, nonadecyl (meth)acrylate, eicosyl (meth)acrylate, cetylcicosyl (meth)acrylate, stearylcicosyl (meth)acrylate, docosyl (meth)acrylate and/or eicosyltetratriacontyl (meth)acrylate;

25

30

35

cycloalkyl (meth)acrylate, such as cyclopentyl (meth)acrylate, cyclohexyl (meth)acrylate, 3-vinyl-2-butylcyclohexyl (meth)acrylate and bornyl (meth)acrylate;

(meth)acrylates which derive from unsaturated alcohols e.g. 2-propynyl (meth)acrylate, allyl (meth)acrylate, and oleyl (meth)acrylate, vinyl (meth)acrylate;

5 The mixtures may, if appropriate, comprise

d) a monomer capable of free-radical polymerization and having at least two terminal olefinic groups whose reactivity differs, for example a
10 bifunctional monomer having a methacrylate end group and a vinyl end group, and/or

e) at least one ethylenically unsaturated monomer (B), preferably from the group of the
15 methacrylates, particularly preferably 2-hydroxyethyl methacrylate.

aminoalkyl (meth)acrylates, such as
tris(2-methacryloxyethyl)amine, N-methylformamidoethyl
20 (meth)acrylate, 2-ureidoethyl (meth)acrylate;

carbonyl-containing (meth)acrylates, such as
2-carboxyethyl (meth)acrylate, carboxymethyl
(meth)acrylate, oxazolidinylethyl (meth)acrylate,
25 N-(methacryloyloxy)formamide, acetonyl (meth)acrylate,
N-methacryloylmorpholine, N-methacryloyl-2-pyrrolidinone;

(meth)acrylates of ether alcohols, e.g.
tetrahydrofurfuryl (meth)acrylate, vinyloxyethoxyethyl
30 (meth)acrylate, methoxyethoxyethyl (meth)acrylate,
1-butoxypropyl (meth)acrylate, 1-methyl-(2-vinyloxy)ethyl
(meth)acrylate, cyclohexyloxymethyl (meth)acrylate,
methoxymethoxyethyl (meth)acrylate, benzyloxymethyl
(meth)acrylate, furfuryl (meth)acrylate, 2-butoxyethyl
35 (meth)acrylate, 2-ethoxyethoxymethyl (meth)acrylate,
2-ethoxyethyl (meth)acrylate, allyloxymethyl
(meth)acrylate, 1-ethoxybutyl (meth)acrylate,
methoxymethyl (meth)acrylate, 1-ethoxyethyl
(meth)acrylate, ethoxymethyl (meth)acrylate;

(meth)acrylates of halogenated alcohols such as
2,3-dibromopropyl (meth)acrylate, 4-bromophenyl
(meth)acrylate, 1,3-dichloro-2-propyl (meth)acrylate,
2-bromoethyl (meth)acrylate, 2-iodoethyl (meth)acrylate,
5 chloromethyl (meth)acrylate;

oxiranyl (meth)acrylates, such as 2,3-epoxybutyl
(meth)acrylate, 3,4-epoxybutyl (meth)acrylate, glycidyl
(meth)acrylate;

10

amides of (meth)acrylic acid, e.g.
N-(3-dimethylaminopropyl) (meth)acrylamide,
N-(diethylphosphono) (meth)acrylamide,
1-(meth)acryloylamido-2-methyl-2-propanol,
15 N-(3-dibutylaminopropyl) (meth)acrylamide,
N-tert-butyl-N-(diethylphosphono) (meth)acrylamide,
N,N-bis(2-diethylaminoethyl) (meth)acrylamide,
4-(meth)acryloylamido-4-methyl-2-pentanol,
N-(methoxymethyl) (meth)acrylamide,
20 N-(2-hydroxyethyl) (meth)acrylamide,
N-acetyl (meth)acrylamide, N-(dimethylaminoethyl)
(meth)acrylamide, N-methyl-N-phenyl (meth)acrylamide,
N,N-diethyl (meth)acrylamide, N-methyl (meth)acrylamide,
N,N-dimethyl (meth)acrylamide, N-isopropyl (meth)acrylamide;

25

heterocyclic (meth)acrylates, such as
2-(1-imidazolyl)ethyl (meth)acrylate,
2-(4-morpholinyl)ethyl (meth)acrylate and
1-(2-methacryloyloxyethyl)-2-pyrrolidone;

30

phosphorus-, boron- and/or silicon-containing
(meth)acrylates, such as
2-(dimethylphosphato)propyl (meth)acrylate,
2-(ethylenephosphito)propyl (meth)acrylate,
35 dimethylphosphinomethyl (meth)acrylate,
dimethylphosphonoethyl (meth)acrylate, diethyl
(meth)acryloylphosphonate, dipropyl (meth)acryloyl
phosphate;

sulphur-containing (meth)acrylates, such as
ethylsulphinyethyl (meth)acrylate, 4-thiocyanatobutyl
(meth)acrylate, ethylsulphonyethyl (meth)acrylate,
thiocyanatomethyl (meth)acrylate, methylsulphinylmethyl
5 (meth)acrylate, bis((meth)acryloyloxyethyl) sulphide;

bis(allyl carbonates), such as ethylene glycol
bis(allyl carbonate), 1,4-butanediol bis(allyl
carbonate), diethylene glycol bis(allyl carbonate);

10 vinyl halides, such as vinyl chloride, vinyl fluoride,
vinylidene chloride and vinylidene fluoride;

vinyl esters, such as vinyl acetate.

15 heterocyclic vinyl compounds, such as 2-vinylpyridine,
3-vinylpyridine, 2-methyl-5-vinylpyridine,
3-ethyl-4-vinylpyridine, 2,3-dimethyl-5-vinylpyridine,
vinylpyrimidine, vinylpiperidine, 9-vinylcarbazole,
20 3-vinylcarbazole, 4-vinylcarbazole, 1-vinylimidazole,
2-methyl-1-vinylimidazole, N-vinylpyrrolidone,
2-vinylpyrrolidone, N-vinylpyrrolidine,
3-vinylpyrrolidine, N-vinylcaprolactam,
N-vinylbutyrolactam, vinyloxolane, vinylfuran,
25 vinylthiophene, vinylthiolane, vinylthiazoles and
hydrogenated vinylthiazoles, vinyloxazoles and
hydrogenated vinyloxazoles;

vinyl ethers and isoprenyl ethers;

30 maleic acid and maleic acid derivatives, such as mono-
and diesters of maleic acid, the alcohol radicals
having from 1 to 9 carbon atoms, maleic anhydride,
methylmaleic anhydride, maleimide, methylmaleimide;

35 fumaric acid and fumaric acid derivatives, such as
mono- and diesters of fumaric acid, the alcohol
radicals having from 1 to 9 carbon atoms.

for completeness, a di(meth)acrylate listed under monomer (A) may also be used as monomer (B).

5 In this context, the term (meth)acrylates encompasses methacrylates and acrylates and also mixtures of the two. Correspondingly, the term (meth)acrylic acid encompasses methacrylic acid and acrylic acid and also mixtures of the two.

10 The ethylenically unsaturated monomers may be used individually or in the form of a mixture.

In principle, the composition of the inventive monomer mixtures may be as desired. It can be utilized to match
15 the property profile of the inventive plastic to the demands of the application. However, it has proven to be highly advantageous to select the composition of the monomer mixture in such a way that the prepolymer composed of the compound(s) of the formula (I), (II)
20 and (III) and the at least one monomer (A) and styrene mix homogeneously at the desired polymerization temperature, because mixtures of this type are easy to handle due to their generally low viscosity and, furthermore, can be polymerized to give homogeneous
25 plastics with better properties.

According to one particularly preferred embodiment of the present invention, the monomer mixture comprises a prepolymer composed of at least 5.0% by weight,
30 preferably at least 20.0% by weight, particularly preferably at least 50.0% by weight, of compounds of the formula (I), (II) and (III), based in each case on the total weight of the monomer mixture. The proportion by weight of the monomer (A) is preferably at least
35 2.0% by weight, preferably at least 10.0% by weight, particularly preferably at least 20.0% by weight, based in each case on the total weight of the monomer mixture. The proportion by weight of aromatic vinyl compounds, in particular styrene, is preferably at

least 2.0% by weight, preferably at least 10.0% by weight, particularly preferably at least 20.0% by weight, based in each case on the total weight of the monomer mixture.

5

According to one particular aspect of the present invention, the mixture comprises
from 50 to 90% by weight, in particular from 60 to 85% by weight, of the prepolymer of the monomers of the
10 formulae (I) and/or (II) and (III),
from 2 to 50% by weight, in particular from 10 to 30% by weight, of monomers (A), and
from 2 to 50% by weight, in particular from 10 to 30% by weight, of aromatic vinyl compounds, in particular
15 styrene, and
from 0 to 45% by weight, in particular from 1 to 10% by weight, of monomers of the formulae (XII) and (XIII),
and/or monomers (B), based in each case on the total weight of the monomer mixture.

20

The preparation of the monomer mixture to be used according to the invention is obvious to the person skilled in the art. By way of example, it can take place via mixing of the prepolymer composed of
25 thio(meth)acrylates of the formulae (I) and/or (II) with (III), and of the aromatic vinyl compounds, and of the monomers (A) and (B), in a manner known per se.

30

For the purposes of the present invention, the monomer mixture is preferably flowable at atmospheric pressure and temperatures in the range from 20.0 to 80.0°C. The term "flowable" is known to the person skilled in the art. It characterizes a liquid which can preferably be cast into various shapes and, using suitable aids,
35 stirred and homogenized. For the purposes of the invention, particular flowable materials have, in particular at 25°C and at atmospheric pressure (101325 Pa) dynamic viscosities of the order of from 0.1 mPa.s to 10 Pa.s, advantageously in the range from

0.65 mPa.s to 1 Pa.s. In one particularly preferred embodiment of the present invention, a cast monomer mixture has no bubbles, in particular no air bubbles. Preference is likewise given to monomer mixtures from
5 which bubbles, in particular air bubbles, can be removed via suitable processes, such as temperature increase and/or application of vacuum.

The inventive high-transparency plastic is obtainable
10 via free-radical copolymerization of the low-viscosity ($\eta < 200$ mPa.s) monomer mixture described above. Free-radical copolymerization is a well-known process initiated via free radicals, converting a mixture of
15 low-molecular monomers into high-molecular-weight compounds, known as polymers. For further details reference is made to the disclosure of H.G. Elias, Makromoleküle [Macromolecules], Volume 1 and 2, Basle, Heidelberg, New York Hüthig und Wepf. 1990 und Ullmann's Encyclopedia of Industrial Chemistry,
20 5th edition, headword "Polymerization Processes".

In one preferred embodiment of the present invention, the inventive plastic is obtainable via mass or bulk polymerization of the monomer mixture. Mass or bulk
25 polymerization here means a polymerization process in which monomers are polymerized without solvent, the polymerization reaction therefore being carried out on the undiluted material or in bulk. Processes which contrast with this are polymerization in emulsion
30 (known as emulsion polymerization) and polymerization in a dispersion (known as suspension polymerization), in which the organic monomers are suspended with protective colloids and/or stabilizers in an aqueous phase, and relatively coarse polymer particles are
35 formed. A particular form of heterogeneous-phase polymerization is bead polymerization, which in essence is a type of suspension polymerization.

In principle, the polymerization reaction may be

initiated in any manner familiar to the person skilled in the art, for example using a free-radical initiator (e.g. peroxide, azo compound) or via irradiation with UV rays or with visible light, α -radiation, β -radiation or γ -radiation, or a combination of these.

In one preferred embodiment of the present invention, lipophilic free-radical polymerization initiators are used to initiate the polymerization. The free-radical polymerization initiators are in particular lipophilic in order to dissolve in the bulk polymerization mixture. Among compounds which may be used, besides the traditional azo initiators, such as azoisobutyronitrile (AIBN) or 1,1-azobiscyclohexanecarbonitrile, are aliphatic peroxy compounds, such as tert-amyl peroxyneodecanoate, tert-amyl peroxy-pivalate, tert-butyl peroxy-pivalate, tert-amyl 2-ethylperoxyhexanoate, tert-butyl 2-ethylperoxyhexanoate, tert-amyl 3,5,5-trimethylperoxyhexanoate, ethyl 3,3-di(tert-amylperoxy)butyrate, tert-butyl perbenzoate, tert-butyl hydroperoxide, decanoyl peroxide, lauryl peroxide, benzoyl peroxide and any desired mixtures of the compounds mentioned. Among the abovementioned compounds, very particular preference is given to AIBN.

In another preferred embodiment of the present invention, the polymerization is initiated by using known photoinitiators, via irradiation with UV rays or the like. Use may be made here of the familiar, commercially available compounds, e.g. benzophenone, α,α -diethoxyacetophenone, 4,4-diethylaminobenzophenone, 2,2-dimethoxy-2-phenylacetophenone, 4-isopropylphenyl 2-hydroxy-2-propyl ketone, 1-hydroxycyclohexyl phenyl ketone, isoamyl p-dimethylaminobenzoate, methyl 4-dimethylaminobenzoate, methyl o-benzoylbenzoate, benzoin, benzoin ethyl ether, benzoin isopropyl ether, benzoin isobutyl ether, 2-hydroxy-2-methyl-1-phenylpropan-1-one, 2-isopropylthioxanthone, dibenzosuberone, 2,4,6-trimethylbenzoyldiphenylphosphine

oxide, bisacylphosphine oxide and other compounds, and the photoinitiators mentioned here may be used alone or in a combination of two or more or in combination with one of the above polymerization initiators.

5

The amount of the free-radical generators may vary widely. By way of example, amounts preferably used are in the range from 0.1 to 5.0% by weight, based on the weight of the entire composition. Particular preference
10 is given to the use of amounts in the range from 0.1 to 2.0% by weight, in particular amounts in the range from 0.1 to 0.5% by weight, based in each case on the weight of the entire composition.

15 The polymerization temperature to be selected for the polymerization is obvious to the person skilled in the art. It is primarily determined via the initiator selected and the manner of initiation (thermal, via irradiation, etc.). It is known that the polymerization
20 temperature can affect the properties of a polymer product. For the purposes of the present invention, preference is therefore given to polymerization temperatures in the range from 20.0 to 100.0°C, advantageously in the range from 20.0 to 80.0°C, in
25 particular in the range from 20.0 to 60.0°C. In one particularly preferred embodiment of the present invention, the reaction temperature is increased during the reaction, preferably in stages.

30 Heat-conditioning at an elevated temperature has also proven to be advantageous, for example from 100 to 150°C, towards the end of the reaction.

The reaction may take place either at subatmospheric
35 pressure or else at superatmospheric pressure. However, it is preferably carried out at atmospheric pressure. The reaction may take place in air or else under an inert gas, preferably with a minimum content of oxygen present, because this content has an inhibiting effect

on any polymerization.

In one particularly preferred embodiment of the present invention, the procedure for preparing the inventive
5 high-transparency plastic prepares a homogeneous mixture from the components, these being monomer mixture, initiator and other additives, e.g. lubricants, and then charges these between glass plates whose shape has been predetermined via the subsequent
10 application, e.g. in the form of spectacle lenses or other lenses, prisms or other optical components. The bulk polymerization is initiated via introduction of energy, for example via high-energy radiation, in particular using UV light, or via heating,
15 advantageously in a water bath and for two or more hours. This gives the optical material in its desired form as clear, transparent, colourless, hard plastic.

For the purposes of the present invention, lubricants
20 are additives for charges of plastic materials, such as compression-moulding materials and injection-moulding materials, their function being to increase the slip capability of the materials charged and thus to ease the moulding of the compression-moulding materials.
25 Examples of substances suitable for this purpose are metal soaps and siloxane combinations. The insolubility of the lubricant in plastics causes some of the lubricant to migrate to the surface during processing, where it acts as a release agent. Particularly suitable
30 lubricants, such as non-ionic fluorinated agents with surface activity, non-ionic silicone agents with surface activity, quaternary alkylammonium salts and acidic phosphate esters, are described in EP 271839 A, the disclosure of which is expressly incorporated by
35 reference for the purposes of the present invention.

The invention provides a high-transparency plastic with very good optical and mechanical properties. For example, its transmittance to DIN 5036 is preferably

greater than 88.0%, advantageously greater than 89.0%.

The refractive index n_D of the inventive plastic is preferably greater than or equal to 1.59. The
5 refractive index of a medium is generally dependent on the wavelength of the incident radiation and on the temperature. The inventive data for refractive index are therefore based on the standard data specified in
10 DIN 53491 (standard wavelength of the (yellow) D line of sodium (about 589 nm)).

According to the invention, the Abbe number of the plastic is preferably > 36.0 to DIN 53491. Information concerning the Abbe number can be found by the person
15 skilled in the art in the literature, for example in the Lexikon der Physik [Dictionary of Physics] (Walter Greulich (ed.); Lexikon der Physik [Dictionary of Physics]; Heidelberg; Spektrum, Akademischer Verlag; Volume 1; 1998).

20 According to one particularly preferred embodiment of the present invention, the plastic has an Abbe number > 36.0 , advantageously > 37.0 , in particular > 38.0 .

25 The FDA falling ball test (ANSI Z 80.1) is used to test mechanical properties. The test is passed if the test specimen is undamaged by a ball of diameter 16 mm. The greater the diameter of the ball used in the test without damaging the specimen, the better the
30 mechanical properties.

The inventive plastic also advantageously has a high glass transition temperature, and therefore maintains its outstanding mechanical properties, in particular
35 its impact strength and its hardness, even at temperatures above room temperature. The glass transition temperature of the inventive plastic is preferably greater than 80°C , advantageously greater than 90°C , in particular greater than 95°C .

Possible fields of use for the inventive high-transparency plastic are obvious to the person skilled in the art. It is particularly suitable for any application destined for transparent plastics. Its
5 characteristic properties make it especially suitable for optical lenses, in particular for ophthalmic lenses.

The following inventive examples and the comparative
10 example serve to illustrate the invention, with no intended resultant restriction.

Examples

15 Synthesis of the thiomethacrylate mixture

75.36 g of 1,2-ethanedithiol are weighed into an Erlenmeyer flask with inert gas feed and stirred, and 416.43 g of 13% strength NaOH solution are metered in
20 within a period of 30 minutes at from 25 to 30°C, with water cooling. A brownish, clear solution forms.

178.64 g of methacrylic anhydride and the Na thiolate solution are then metered in parallel at the desired
25 metering temperature within a period of 45 minutes into the initial charge of stirred ethyl acetate/water in the reaction flask. Where appropriate here, inert gas is passed over the mixture. The contents of the flask generally become cooler by about 2°C at the start of
30 the feed, and a slightly exothermic reaction begins after about 5-10 minutes, meaning that appropriate cooling is applied in order to maintain the desired reaction temperature (35°C). Once the feed has ended, the mixture is stirred for a further 5 minutes at 35°C
35 and is then cooled, with stirring, to about 25°C.

The mixture is transferred to a separating funnel and separated, and the lower, aqueous phase is discharged. For work-up, the organic phase is transferred to an

Erlenmeyer flask and stirred with ®Dowex M-31 for about 15 minutes, the ion exchanger then being filtered off.

5 The somewhat cloudy to almost clear crude ester solution is then stabilized with 100 ppm of HQME and concentrated at not more than 50°C on a rotary evaporator. The colourless final product is filtered at room temperature (20-25°C). This gives about 140 g of colourless, clear ester.

10

Preparation of prepolymer: reaction of 6.84 g of the thiodi(meth)acrylate and 0.36 g of DMDO in the presence of an amine as catalyst, the method being based on EP 284374.

15

In an example of the preparation of a polymer based on an oligomeric thiodimethacrylate, 7.2 g of the prepolymer, 2.4 g of styrene, 2.4 g of decaethoxylated bisphenol A di(meth)acrylate, 0.1 g of hydroxyethyl
20 methacrylate, 36 mg of a UV initiator, e.g. Irgacur 819, and 24 mg of tert-butyl peroctoate or similar initiators (cf. Inventive Example 1) are mixed. The homogeneous casting resin mixture is placed in an appropriate mould and hardened within a period of 10
25 min in a UV curing system using a 1200 W high-pressure mercury source. The material is then heat-conditioned for a further period of about 2 h at about 120°C in an oven.

Experiment	System	Refractive/Abbe coefficient DIN 53491	Odour	Trans- mittance	FDA falling ball test diameter ANSI Z80.1	Average ball diameter in mm, test passed
Inventive Examples IE 1	PLEX 6931/DMDO prepolymer co styrene co E10BADMA = 60:20:20 plus 1% HEMA	589 nm 1.5939	no	89	passed	18
Comparative Examples						
CE I	PLEX 6931 co DMDO co styrene co E10BADMA = 57:3:20:20 plus 1% HEMA (#)	-	yes	-	-	-
CE II	PLEX 6931 co styrene co E10BADMA = 60:20:20 plus 1% HEMA	1.5959	no	89	passed	16
CE III	PLEX 6931/DMDO prepolymer co styrene = 70:30 plus 1% HEMA	1.6089	no	89	passed	16

Plex 6931 O: reaction product from methacrylic anhydride and ethanedithiol from DE 316671

E10BADMA: ethoxylated bisphenol A dimethacrylate, degree of ethoxylation about 10

DMDO: dimercaptodioxaoctane

HEMA: hydroxyethylmethacrylate

(#): no further analysis was undertaken, because of the problems of odour.

The inventive mixture (IE 1) is odourless. The Comparative Example CE I did not pass this test and was therefore not studied further.

- 5 For comparable refractive index (of IE 1 with CE II and CE III) the Abbe number was nevertheless better for the inventive mixture. The inventive mixture also performed substantially better in the falling ball test.